

50618204 8/12/06

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* * * * * Welcome to STN International * * * * *

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America
NEWS 2 "Ask CAS" for self-help around the clock
NEWS 3 DEC 21 IPC search and display fields enhanced in CA/CAplus with the
IPC reform
NEWS 4 DEC 23 New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
USPAT2
NEWS 5 JAN 13 IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS 6 JAN 13 New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
INPADOC
NEWS 7 JAN 17 Pre-1988 INPI data added to MARPAT
NEWS 8 JAN 17 IPC 8 in the WPI family of databases including WPIFV
NEWS 9 JAN 30 Saved answer limit increased
NEWS 10 JAN 31 Monthly current-awareness alert (SDI) frequency
added to TULSA
NEWS 11 FEB 21 STN AnaVist, Version 1.1, lets you share your STN AnaVist
visualization results
NEWS 12 FEB 22 Status of current WO (PCT) information on STN
NEWS 13 FEB 22 The IPC thesaurus added to additional patent databases on STN
NEWS 14 FEB 22 Updates in EPFULL; IPC 8 enhancements added
NEWS 15 FEB 27 New STN AnaVist pricing effective March 1, 2006
NEWS 16 FEB 28 MEDLINE/LMEDLINE reload improves functionality
NEWS 17 FEB 28 TOXCENTER reloaded with enhancements
NEWS 18 FEB 28 REGISTRY/ZREGISTRY enhanced with more experimental spectral
property data
NEWS 19 MAR 01 INSPEC reloaded and enhanced
NEWS 20 MAR 03 Updates in PATDPA; addition of IPC 8 data without attributes
NEWS 21 MAR 08 X.25 communication option no longer available after June 2006
NEWS 22 MAR 22 EMBASE is now updated on a daily basis

NEWS EXPRESS FEBRUARY 15 CURRENT VERSION FOR WINDOWS IS V8.01a,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
V8.0 AND V8.01 USERS CAN OBTAIN THE UPGRADE TO V8.01a AT
<http://download.cas.org/express/v8.0-Discover/>

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 11:21:11 ON 01 APR 2006

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 11:21:19 ON 01 APR 2006

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 30 MAR 2006 HIGHEST RN 878649-01-5

DICTIONARY FILE UPDATES: 30 MAR 2006 HIGHEST RN 878649-01-5

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

*
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *
*

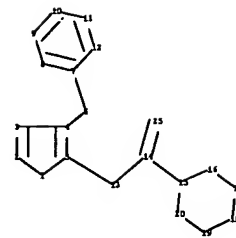
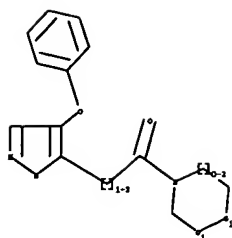
Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>

Uploading C:\Program Files\Stnexp\Queries\10669794.str



chain nodes :
6 13 14 25
ring nodes :
1 2 3 4 5 7 8 9 10 11 12 15 16 17 18 19 20
chain bonds :
4-6 5-13 6-7 13-14 14-15 14-25
ring bonds :
1-2 1-5 2-3 3-4 4-5 7-8 7-12 8-9 9-10 10-11 11-12 15-16 15-20 16-17
17-18 18-19 19-20
exact/norm bonds :
1-2 1-5 2-3 3-4 4-5 4-6 5-13 6-7 13-14 14-15 14-25 15-16 15-20 16-17
17-18 18-19 19-20
normalized bonds :
7-8 7-12 8-9 9-10 10-11 11-12
isolated ring systems :
containing 1 : 7 :

G1:C,N

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:Atom 8:Atom 9:Atom 10:Atom
11:Atom 12:Atom 13:CLASS 14:CLASS 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom
20:Atom 25:CLASS

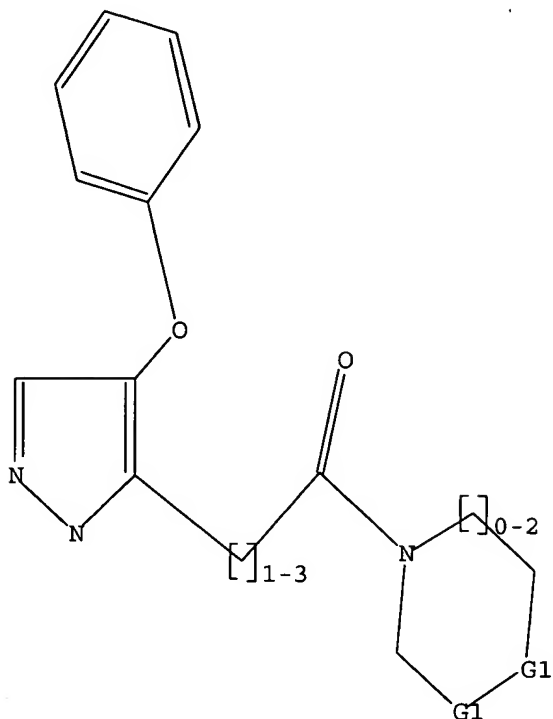
50618204 8/12/06

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



G1 C,N

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 11:21:42 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS

SEARCH TIME: 00.00.01

0 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

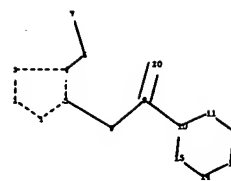
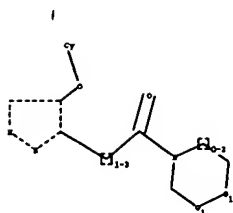
PROJECTED ITERATIONS: 0 TO 0

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=>

Uploading C:\Program Files\Stnexp\Queries\106697941.str



chain nodes :
6 7 8 9 20
ring nodes :
1 2 3 4 5 10 11 12 13 14 15
chain bonds :
4-6 5-8 6-7 8-9 9-10 9-20
ring bonds :
1-2 1-5 2-3 3-4 4-5 10-11 10-15 11-12 12-13 13-14 14-15
exact/norm bonds :
1-2 1-5 2-3 3-4 4-5 4-6 5-8 6-7 8-9 9-10 9-20 10-11 10-15 11-12 12-13
13-14 14-15
isolated ring systems :
containing 1 : 7 :

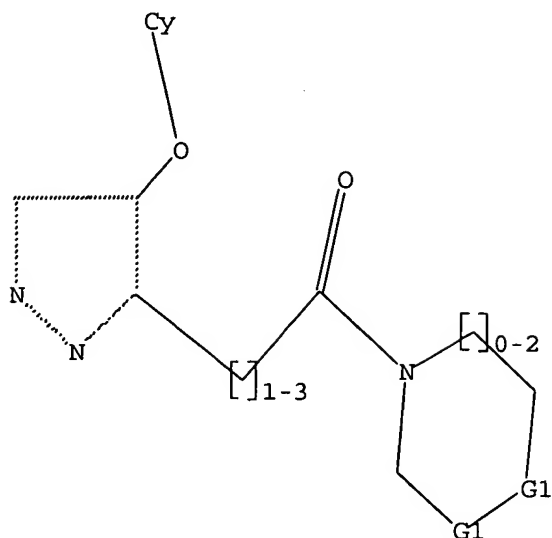
G1:C,N

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:Atom 8:CLASS 9:CLASS 10:Atom
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 20:CLASS

L3 STRUCTURE UPLOADED

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=> d
L3 HAS NO ANSWERS
L3 STR



G1 C,N

Structure attributes must be viewed using STN Express query preparation.

=> s l3
SAMPLE SEARCH INITIATED 11:26:35 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 0 TO 0
PROJECTED ANSWERS: 0 TO 0

L4 0 SEA SSS SAM L3

=> s l4 full
FULL SEARCH INITIATED 11:28:53 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 23 TO ITERATE

100.0% PROCESSED 23 ITERATIONS 20 ANSWERS
SEARCH TIME: 00.00.01

L5 20 SEA SSS FUL L3

=> file caplus
COST IN U.S. DOLLARS
FULL ESTIMATED COST
SINCE FILE ENTRY 172.22
TOTAL SESSION 172.43

FILE 'CAPLUS' ENTERED AT 11:28:59 ON 01 APR 2006

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FILE COVERS 1907 - 1 Apr 2006 VOL 144 ISS 15
FILE LAST UPDATED: 31 Mar 2006 (20060331/ED)

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=> s 15

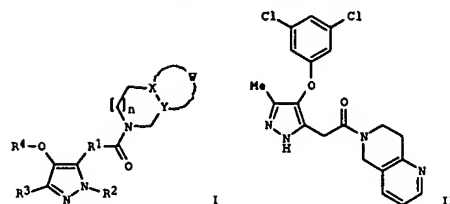
L6 2 L5

=> d ibib abs hitstr tot

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:292024 CAPLUS
 DOCUMENT NUMBER: 140:303665
 TITLE: Preparation of pyrazole amides for treating HIV infections
 INVENTOR(S): Jones, Lyn Howard; Mowbray, Charles Eric; Price, David
 Anthony; Selby, Matthew Duncan; Stupples, Paul Anthony
 PATENT ASSIGNEE(S): Pfizer Limited, UK; Pfizer Inc.
 SOURCE: PCT Int. Appl., 55 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004029051	A1	20040408	WO 2003-1B4071	20030915
V: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GR, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2495338	AA	20040408	CA 2003-2495338	20030915
AU 2003263455	A1	20040409	AU 2003-263455	20030915
BR 2003014759	A	20050726	BR 2003-14759	20030915
EP 1556381	A1	20050727	EP 2003-798295	20030915
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2006505625	T2	20060216	JP 2005-501939	20030915
US 2005004129	A1	20050106	US 2003-669794	20030923
PRIORITY APPLN. INFO.:				
GB 2002-22375 A 20020926				
GB 2002-23357 A 20021008				
US 2002-433220P P 20021213				
WO 2003-1B4071 W 20030915				
OTHER SOURCE(S): MARPAT 140:303665				
GI				

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



AB The title compds. [I: WXY = (un)substituted 5-6 membered partially saturated or aromatic ring containing 0-3 N atoms wherein X = CH or N and Y = CH, or, when X = CH, may also be N; R1 = a bond, alkylene, R2 = H, alkyl, cycloalkyl, etc.; R3 = H, alkyl, cycloalkyl, etc.; R4 = (un)substituted Ph, naphthyl, pyridyl; n = 0-2] which bind to the enzyme reverse transcriptase and are modulators, especially inhibitors thereof, were prepared and formulated.

Thus, reacting [4-(3,5-dichlorophenoxy)-3-methyl-1H-pyrazol-5-yl]acetic acid (preparation given) with 5,6,7,8-tetrahydro-[1,6]naphthyridine afforded II. The compds. I were tested for inhibition of HIV-1 reverse transcriptase enzyme (data were given for representative compds. I). The compds. I are useful in the treatment of a variety of disorders including those in which the inhibition of reverse transcriptase is implicated. Disorders of interest include those caused by Human Immunodeficiency Virus (HIV) and genetically related retroviruses, such as Acquired Immune Deficiency Syndrome (AIDS).

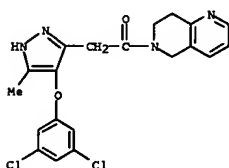
IT 676994-33-SP 676994-34-6P 676994-35-7P
 676994-36-8P 676994-38-0P 676994-39-1P
 676994-40-4P 676994-41-5P 676994-42-6P
 676994-43-7P 676994-44-8P 676994-45-9P
 676994-47-1P 676994-48-2P 676994-49-3P
 676994-50-6P 676994-51-7P 676994-52-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PRFP (Preparation); USES (Uses)

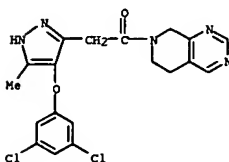
(preparation of pyrazole amides for treating HIV infections)

RN 676994-33-5 CAPLUS
 CN 1,6-Naphthyridine, 6-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

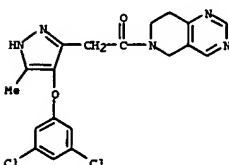
L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 676994-34-6 CAPLUS
 CN Pyrido[3,4-d]pyrimidine, 7-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

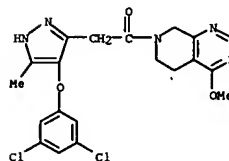


RN 676994-35-7 CAPLUS
 CN Pyrido[4,3-d]pyrimidine, 6-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

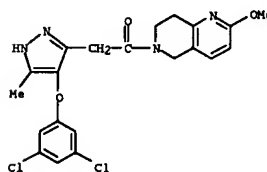


RN 676994-36-8 CAPLUS
 CN Pyrido[3,4-d]pyrimidine, 7-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro-4-methoxy- (9CI) (CA INDEX NAME)

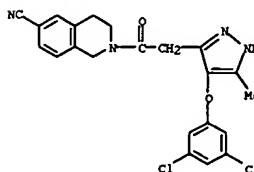
L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 676994-38-0 CAPLUS
 CN 1,6-Naphthyridine, 6-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro-2-methoxy- (9CI) (CA INDEX NAME)

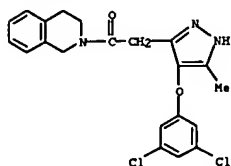


RN 676994-39-1 CAPLUS
 CN 6-Isquinolinescarbonitrile, 2-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-1,2,3,4-tetrahydro- (9CI) (CA INDEX NAME)

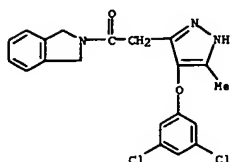


RN 676994-40-4 CAPLUS
 CN 6-Isquinolinescarbonitrile, 2-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-1,2,3,4-tetrahydro- (9CI) (CA INDEX NAME)

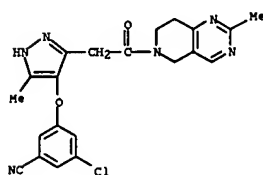
L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 676994-41-5 CAPLUS
CN 1H-isoindole, 2-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-2,3-dihydro- (9CI) (CA INDEX NAME)

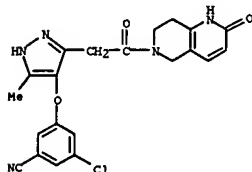


RN 676994-42-6 CAPLUS
CN Pyrido[4,3-d]pyrimidine, 6-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro-2-methyl- (9CI) (CA INDEX NAME)

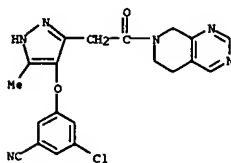


RN 676994-43-7 CAPLUS
CN 1,6-naphthyridine, 6-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro-2-(trifluoromethyl)- (9CI) (CA INDEX NAME)

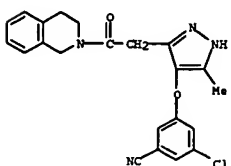
L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 676994-48-2 CAPLUS
CN Pyrido[3,4-d]pyrimidine, 7-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

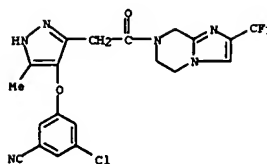


RN 676994-49-3 CAPLUS
CN Isoquinoline, 2-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-1,2,3,4-tetrahydro- (9CI) (CA INDEX NAME)

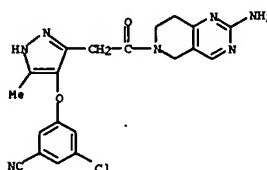


RN 676994-50-6 CAPLUS
CN Pyrido[4,3-d]pyrimidine, 6-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

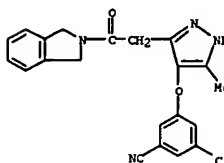
L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 676994-44-8 CAPLUS
CN Pyrido[4,3-d]pyrimidin-2-amine, 6-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

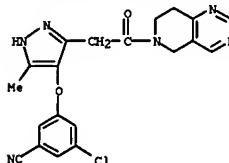


RN 676994-45-9 CAPLUS
CN 1H-isoindole, 2-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-2,3-dihydro- (9CI) (CA INDEX NAME)

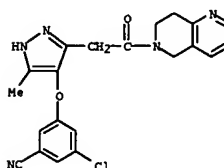


RN 676994-47-1 CAPLUS
CN 1,6-naphthyridine, 6-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)

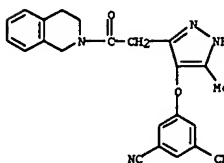
L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 676994-51-7 CAPLUS
CN 1,6-naphthyridine, 6-[[4-(3-chloro-5-cyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-5,6,7,8-tetrahydro- (9CI) (CA INDEX NAME)



RN 676994-52-8 CAPLUS
CN Isoquinoline, 2-[[4-(3,5-dicyanophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-1,2,3,4-tetrahydro- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

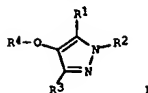
L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2002:832763 CAPLUS
 DOCUMENT NUMBER: 137:337884
 TITLE: Preparation of aryloxy pyrazole derivatives as reverse transcriptase inhibitors for treating HIV
 INVENTOR(S): Jones, Lyn Howard; Mowbray, Charles Eric; Price, Davis Anthony; Selby, Matthew Duncan; Stupples, Paul Anthony
 PATENT ASSIGNEE(S): Pfizer Limited, UK; Pfizer Inc.
 SOURCE: PCT Int. Appl., 306 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002085860	A1	20021031	WO 2002-181234	20020404
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, GR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2443449	AA	20021031	CA 2002-244349	20020404
EP 1377556	A1	20040107	EP 2002-708600	20020404
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
EE 200300497	A	20040216	EE 2003-497	20020404
BR 2002008811	A	20040309	BR 2002-8811	20020404
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JP 2004531535	T2	20041014	JP 2002-583387	20020404
NZ 529403	A	20050624	NZ 2002-529403	20020404
US 2003100554	A1	20030529	US 2002-118512	20020405
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NO 2003004523	A	20031209	NO 2003-4523	20031009
US 2006020012	A1	20060126	US 2005-157340	20050620

PRIORITY APPLN. INFO.:

OTHER SOURCE(S): MARPAT 137:337884
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L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)

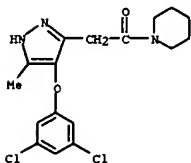


AB This invention relates to pyrazole derivs. (shown as I; e.g. 2-Amino-6-[[4-(3,5-dichlorophenoxy)-3,5-diethyl-1H-pyrazol-1-yl]methyl]-4(3H)-pyrimidinone) or pharmaceutically acceptable salts, solvates or derivative thereof, wherein R1 to R4 are defined below, and to processes for the preparation thereof, intermediates used in their preparation of, compds. containing

them and the uses of such derivs. The compds. of the present invention bind to the enzyme reverse transcriptase and are modulators, especially inhibitors thereof. As such the compds. of the present invention are useful in the treatment of a variety of disorders including those in which the inhibition of reverse transcriptase is implicated. Disorders of interest include those caused by Human Immunodeficiency Virus (HIV) and genetically related retroviruses, such as Acquired Immune Deficiency Syndrome (AIDS). In tests of inhibition of HIV-1 reverse transcriptase enzyme, the claimed compds. 2-amino-6-[[4-(3,5-dichlorophenoxy)-3,5-diethyl-1H-pyrazol-1-yl]methyl]-4(3H)-pyrimidinone, 3,5-dimethyl-4-[[3,5-diethyl-1-(2-hydroxyethyl)-1H-pyrazol-4-yl]oxy]benzonitrile and 1-(3-azetidinyl)-4-(3,5-dichlorophenoxy)-3,5-diethyl-1H-pyrazole had IC50 values of 39,000, 3,200 and 248 nM, resp. In I: R1 is H, C1-C6 alkyl, C3-C7 cycloalkyl, Ph, benzyl, halo, -CN, -OR7, -CO2R10, -CONR5R10, R8 or R9. R2 is H, C1-C6 alkyl, C3-C6 alkenyl, C3-C6 alkynyl, C3-C7 cycloalkyl, C3-C7 cycloalkenyl, Ph, benzyl, R8 or R9; or, R1 and R2, when taken together, represent unbranched C3-C4 alkylene. R3 is H, C1-C6 alkyl, C3-C7 cycloalkyl, Ph, benzyl, halo, -CN, -OR7, -CO2R5, -CONR5R5, R8 or R9; R4 is Ph, naphthyl or pyridyl. Definitions of R5 and R7-R10 and addnl. specifications are given in the claims. Included are 283 claimed-compound preps. and 115 intermediate preps.

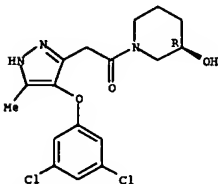
IT 473922-03-1P, 1-[[4-(3,5-Dichlorophenoxy)-3-methyl-1H-pyrazol-5-yl]acetyl]piperidine 473922-04-2P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (Drug candidate; preparation of aryloxy pyrazole derivs. as reverse transcriptase inhibitors for treating HIV)
 RN 473922-03-1 CAPLUS
 CN Piperidine, 1-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]- (9CI) (CA INDEX NAME)

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN (Continued)



RN 473922-04-2 CAPLUS
 CN 3-piperidinol, 1-[[4-(3,5-dichlorophenoxy)-5-methyl-1H-pyrazol-3-yl]acetyl]-, (3R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

10.68

183.11

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

CA SUBSCRIBER PRICE

-1.50

-1.50

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